

**Amendments to the Claims:**

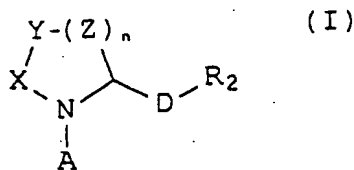
This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1-72 (Canceled)

73. (Currently Amended) A method of treating a neurological disorder selected from the group consisting of peripheral neuropathies caused by physical injury or disease state, physical damage to the brain, physical damage to the spinal cord, stroke associated with brain damage, and neurological disorders relating to neurodegeneration in an animal, comprising:

administering to the animal an effective amount of a compound to stimulate growth of damaged peripheral nerves or to promote neuronal regeneration, where the compound has the formula (I):

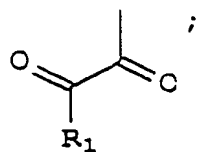


where

X, Y, and Z are independently selected from the group consisting of C, O, S, or N ,  
provided that X, Y, and Z are not all C;

n is 1;

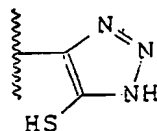
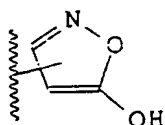
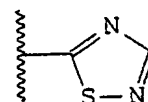
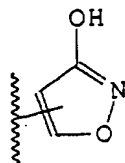
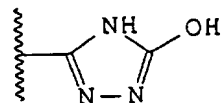
A is



R<sub>1</sub> is selected from the group consisting of hydrogen, C<sub>1</sub>-C<sub>9</sub> straight or branched chain alkyl, C<sub>2</sub>-C<sub>9</sub> straight or branched chain alkenyl, aryl, heteroaryl, carbocycle, and heterocycle;

D is selected from the group consisting of a bond, C<sub>1</sub>-C<sub>10</sub> straight or branched chain alkylene, ethylene (-CH=CH-), and butylene;

R<sub>2</sub> is a carboxylic acid or selected from the group consisting of:



wherein said alkyl, alkenyl, alkylene, ethylene, butylene, aryl, heteroaryl, carbocycle, heterocycle, or  $R_2$  is optionally substituted with one or more substituents selected from  $R_3$ , where

$R_3$  is selected from the group consisting of hydrogen, hydroxy, halo, haloalkyl, thiocarbonyl, alkoxy, alkenoxy, alkylaryloxy, aryloxy, arylalkyloxy, cyano, nitro, imino, alkylamino, aminoalkyl, sulfhydryl, thioalkyl, alkylthio, sulfonyl,  $C_1$ - $C_6$  straight or branched chain alkyl,  $C_2$ - $C_6$  straight or branched chain alkenyl or alkynyl, aryl, heteroaryl, carbocycle, heterocycle, and  $CO_2R_4$  where  $R_4$  is selected from the group consisting of hydrogen,  $C_1$ - $C_9$  straight or branched chain alkyl, and  $C_2$ - $C_9$  straight or branched chain alkenyl;

or a pharmaceutically acceptable salt, or solvate thereof.

74. (Canceled)

75. (Original) The method of claim 73, wherein the neurological disorder is selected from the group consisting of Alzheimer's Disease, Parkinson's Disease, and amyotrophic lateral sclerosis.

76. (Original) The method of claim 73, wherein the neurological disorder is Alzheimer's disease.

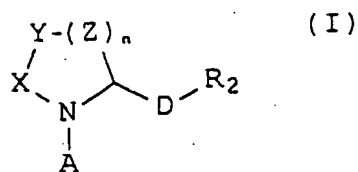
77. (Original) The method of claim 73, wherein the neurological disorder is amyotrophic lateral sclerosis.

78. (Original) The method of claim 73, wherein said compound is non-immunosuppressive.

79. (Previously Presented) The method of claim 73, wherein Y is O, S, or N; R<sub>1</sub> is C<sub>1</sub>-C<sub>9</sub> straight or branched chain alkyl or aryl; and D is a bond or CH<sub>2</sub>.

80. (Previously Presented) A method of treating a neurological disorder in an animal, comprising:

administering to the animal an effective amount of a compound to stimulate growth of damaged peripheral nerves or to promote neuronal regeneration, wherein the compound has the formula (I):

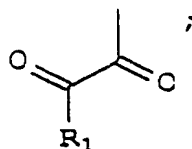


where

X, Y, and Z are independently selected from the group consisting of C, O, S, or N, provided that X, Y, and Z are not all C;

n is 1;

A is



R<sub>1</sub> is selected from the group consisting of hydrogen, C<sub>1</sub>-C<sub>9</sub> straight or branched chain alkyl, C<sub>2</sub>-C<sub>9</sub> straight or branched chain alkenyl, aryl, heteroaryl, carbocycle, and heterocycle;

D is selected from the group consisting of a bond, C<sub>1</sub>-C<sub>10</sub> straight or branched chain alkylene, ethylene (-CH=CH-), and butylene;

R<sub>2</sub> is selected from the group consisting of:

-COOH, -SO<sub>3</sub>H, -SO<sub>2</sub>HNR<sub>3</sub>, -PO<sub>2</sub>H, -CN, -PO(OH)(OR<sub>3</sub>), -C(O)NHOH, -C(O)NHSO<sub>2</sub>R<sub>3</sub>, and -CONHCN;

wherein said alkyl, alkenyl, alkylene, ethylene, butylene, aryl, heteroaryl, carbocycle, heterocycle, or  $R_2$  is optionally substituted with one or more substituents selected from  $R_3$ , where

$R_3$  is selected from the group consisting of hydrogen, hydroxy, halo, haloalkyl, thiocarbonyl, alkoxy, alkenoxy, alkylaryloxy, aryloxy, arylalkyloxy, cyano, nitro, imino, alkylamino, aminoalkyl, sulfhydryl, thioalkyl, alkylthio, sulfonyl,  $C_1$ - $C_6$  or branched chain alkyl,  $C_2$ - $C_6$  straight or branched chain alkenyl or alkynyl, aryl, heteroaryl, carbocycle, heterocycle, and  $CO_2R_4$  where  $R_4$  is selected from the group consisting of hydrogen,  $C_1$ - $C_9$  straight or branched chain alkyl, and  $C_2$ - $C_9$  straight or branched chain alkenyl;

or a pharmaceutically acceptable salt, or solvate thereof.

81. (Previously Presented) The method of claim 80, wherein the neurological disorder is selected from the group consisting of peripheral neuropathies caused by physical injury or disease state, physical damage to the brain, physical damage to the spinal cord, stroke associated with brain damage, and neurological disorders relating to neurodegeneration.

82. (Previously Presented) The method of claim 80, wherein the neurological disorder is selected from the group consisting of Alzheimer's Disease, Parkinson's Disease, and amyotrophic lateral sclerosis.

83. (Previously Presented) The method of claim 80, wherein the neurological disorder is Alzheimer's Disease.

84. (Previously Presented) The method of claim 80, wherein the neurological disorder is amyotrophic lateral sclerosis.

85. (Previously Presented) The method of claim 80, wherein said compound is non-immunosuppressive.

86. (Previously Presented) The method of claim 80, wherein the compound is selected from the group consisting of:

